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(54) Title: CARBOXAMIDE DERIVATIVES AS SELECTIVE INHIBITORS OF PATHOGENS

(57) Abstract: The present invention relates to the use of carboxamide compounds as selective inhibitors of pathogens, particularly viruses and, more particularly, herpesviridae. Surprisingly, these compounds show reduced side effects in comparison with previous antiviral compounds. Thus, a novel method for preventing or treating infections by pathogens, particularly herpesviridae is provided.

Carboxamide derivatives as selective inhibitors of pathogens**Description**

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The present invention relates to the use of carboxamide compounds as selective inhibitors of pathogens, particularly viruses and, more particularly, herpesviridae. Surprisingly, these compounds show reduced side effects in comparison with previous antiviral compounds. Thus, a novel method for preventing or treating infections by pathogens, particularly herpesviridae is provided.

Human Cytomegalovirus (HCMV) is a highly species specific β -herpesvirus. Primary infection of healthy children and adults is usually asymptomatic, with a minority of cases developing a mononucleosis-like syndrome. In contrast, congenital infection (U.S. 0.2%-2.2% per live birth; approx. 40,000 per year) leads to several neurological defects in 10 to 15% of infected neonates. Immunocompromised patients are another group of hosts facing serious diseases caused by HCMV infection or reactivation of a persistent infection. Up to 40% of AIDS patients, for example, develop retinitis, pneumonitis, gastroenteritis or disseminated HCMV disease. Allograft recipients (20,000 allograft transplantations per year in US) are often infected (or superinfected) by virus from the transplanted organ. Clinical symptoms in the postransplant period include prolonged fever, leukopenia, thrombocytopenia, atypical lymphocytosis, elevated hepatic transaminases and decreased graft survival. In bone marrow transplantations, HCMV infection has been associated with high mortality rates (80-90% for untreated HCMV pneumonia), which have been reduced by newer antiviral agents to 10-20% (reviewed in Britt J.B. and Alford C.A., 1996, Cytomegaloviruses, pp.2494-2523. In B.N.Fields, D.M.Knipe, P.M.Howley (ed.) Fields Virology, Lippencott-Raven Pub., Philadelphia).

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Ganciclovir is available for intravenous and oral administration and as an implant in the case of retinitis. Toxicities include leukopenia and thrombocytopenia. Foscarnet (phosphonoformic acid) exhibits considerable renal toxicity and is only available in intravenous form, which is also true for 5 Cidofovir. Although treatment of HCMV-induced disease has been substantially improved with these inhibitors of the viral DNA polymerase and preemptive or early antiviral therapy in transplant patients (Hebart, H. et al., Drugs, 1998, 55:59-72), there is room for improvement in the toxicity profile. Especially in the treatment of retinitis in AIDS patients, where CMV 10 infection has to be controlled for long periods of time and replication will resume once Ganciclovir and Foscarnet are removed, new classes of drugs are needed with better oral bioavailability and activity against emerging Ganciclovir-and Foscarnet-resistant strains.

15 Leflunomide, an immunomodulatory drug used in rheumatoid arthritis, was previously found to inhibit HCMV replication in cell culture. The antiinflammatory and immunosuppressive properties of Leflunomide have been demonstrated in animal models of autoimmune disease and organ transplant rejection (Reviewed in J.Rheumatology (1998) 25:20, Agents Actions (1991) 32:10) and it was recently approved for use in rheumatoid 20 arthritis in the US (Scrip (1998) (2370):22). The proposed basis for the antiproliferative action of Leflunomide, for example, in mitogen-stimulated human T-lymphocytes (JBC (1998) 273:21682), is the suppression of the de novo pathway of pyrimidine synthesis. The active metabolite of 25 Leflunomide is a noncompetitive inhibitor of Dihydroorotate dehydrogenase (DHODH) (Biochem. Pharmacol. (1995) 50:861; J.Pharmacol.Exper.ther.(1995) 275:1043; JBC (1995) 270:22467, Eur.J.Biochem.(1996) 240:292; Biochemistry (1996) 35:1270), a mitochondrial enzyme which catalyzes the rate-limiting step in this 30 biosynthetic pathway. Most research groups favor the inhibition of DHODH as the major mechanism for Leflunomide-induced cytostatic effects (cited in JBC 273, 21682), however, inhibition of tyrosine kinases by Leflunomide

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has been demonstrated in animal models and proposed to be a second important mode of action in immunosuppression (J.Immunol.(1997) 159:22; JBC (1995) 270:12398; Biochem.Pharmacol.(1996) 52:527, J.Immunol.159:167). Tyrosine kinase inhibition is not affected by the 5 addition of uridine, while DHODH inhibition can be reversed by uridine. In the case of Brequinar Sodium, which is another inhibitor of DHODH and an antimetabolite tested for cancer therapy (Cancer Res.(1986) 46:5014, Cancer Chemother.Pharmacol.(1993) 32:64), the finding that only 10 antiproliferative effects were reversed by uridine led to a patent claiming the reduction of non-immunosuppressive side effects by coadministration of uridine (Williams J.W., Chong, A., Xu, X.: WO 98/13047 (1994)).

Surprisingly, it was found that carboxamide derivatives can selectively 15 inhibit kinases from pathogens, particularly from herpesviridae such as HCMV, without significantly inhibiting DHODH. Particularly, these compounds show a pronounced reduction in HCMV replication but do not inhibit purified recombinant human DHODH in an in vitro assay. This 20 surprising uncoupling of antipathogen efficacy and inhibition of the de novo pyrimidine synthesis suggests that the subject carboxamide derivatives are free of the side effects associated with the antiproliferative and immunosuppressive properties of prior art medicaments such as Leflunomide.

These compounds have been shown previously to inhibit platelet-derived 25 growth factor receptor activity (WO 95/19169 Sugen, Inc.), to have antibacterial activity (US 3,303,201) and to inhibit HIV reverse transcriptase (WO 96/16675 Rega Institut). A selective activity of these compounds against kinases from pathogens such as bacteria, protozoa or viruses, particularly without undesired side effects, has not yet been described.

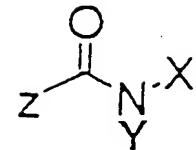
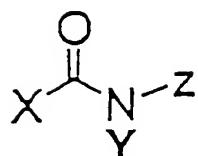
30 Thus, the present invention refers to the use of compounds of the general formulae (Ia), (Ib), (Ic) or (Id):

- 4 -

(Ia)

(Ib)

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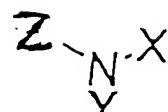
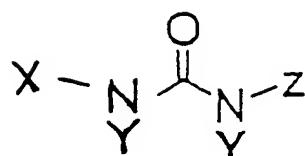


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(Ic)

(Id)

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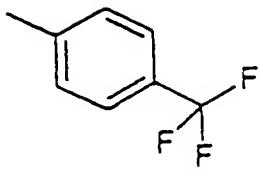
wherein X and Z are substituents comprising an aromatic or heteroaromatic ring system for the manufacture of an agent against infectious diseases. For example, X and/or Z may be an aromatic radical, preferably a phenyl radical which is unsubstituted or which carries at least one substituent, e.g. 1-4 substituents which may be selected from hydroxy, cyano, nitro, halo, e.g. F, Cl, Br, I, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkoxy, C₁-C₄ alkylthio, C₁-C₄ haloalkylthio, carboxy, carboxy-C₁-C₄-alkyl, carboxy-aryl (or heteroaryl), amino carbonyl-C₁-C₄ alkyl, amino carbonyl - aryl (or heteroaryl) and aryl (heteroaryl), Y is hydrogen or C₁-C₄ alkyl.

Especially preferred are compounds, wherein X and/or Z is an aromatic or heteroaromatic radical, e.g. a phenyl radical having at least one, particularly one, two or three C₁-C₄ haloalkyl substituents, e.g. C₁ haloalkyl substituents such as -CF₃, -CHF₂ and -CH₂F.

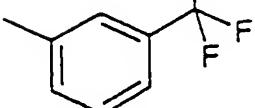
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For example, X and/or Z may be selected from radicals represented by formulae (IIa-c):

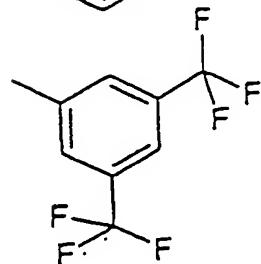
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20 Y is preferably hydrogen or C₁-C₂ alkyl, more preferably hydrogen.

In a further preferred embodiment, X and/or Z comprises a heterocyclic ring which may contain one or several heteroatoms such as oxygen, nitrogen and/or sulfur, preferably a 5-membered heterocyclic ring which may be selected from pyrrole, pyrazole, imidazole, 1,2,3-triazole, tetrazole, oxazole, isoxazole, thiazole, isothiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole, thiophene, furan, indole and 3-thiaindole, a 6-membered heterocyclic ring which may be selected from pyridine, pyran, pyrimidine, pyridazine, pyrimidine and pyrazine, or a bi- or polycyclic heteroaromatic ring such as indazole, imidazole, chinoline or isochinoline. The heterocyclic ring can be mono, di, tri or tetrasubstituted with substituents as defined above.

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For example, X and/or Z may be selected from radicals represented by formulae (IIIa-k):

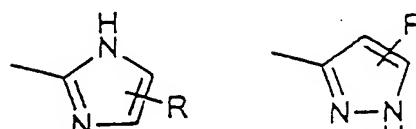
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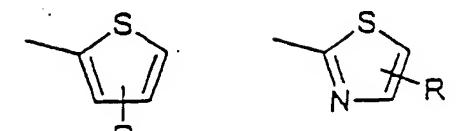
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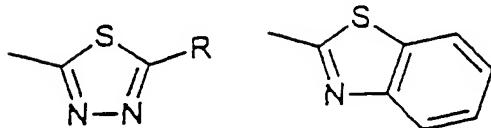
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wherein R is at least one substituent, e.g. one or two substituents selected from halo, C₁-C₃ alkyl, C₁-C₃ alkoxy or aryl, e.g. phenyl.

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The aromatic and/or heteroaromatic ring of X and Z may be directly linked to the central structural element of the compounds (Ia-d), i.e. by a covalent

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bond. Alternatively, the linkage may comprise an alkylene-group, preferably a C₁-C₄ alkylene group, e.g. a C₁ or C₂ alkylene group.

Surprisingly, compounds (Ia), (Ib), (Ic) and (Id) have a selectivity for kinases

5 from pathogens compared with the cellular enzyme DHODH. Thus, the compounds are suitable for the preparation of an agent against infectious diseases, particularly viral infections, more particularly, against infections

by herpesviruses. The herpesviruses may be selected from human herpesviruses and herpesviruses from other mammals, such as bovine,

10 equine, porcine and pongine herpesviruses. Suitable herpesviruses are selected from α -herpesviruses, e.g. simplexviruses such as herpes simplex virus 1, herpes simplex virus 2, bovine herpesvirus 2, cercopithecine

herpesvirus 1 or varicellaviruses such as varicella zoster virus, porcine herpesvirus 1 (pseudorabiesvirus) bovine herpesvirus 1 and equine

15 herpesvirus 1 (equine abortion virus). Further, the herpesvirus may be selected from β -herpesviruses, e.g. cytomegaloviruses such as human cytomegalovirus and from roseoloviruses, such as human herpesvirus 6,

human herpesvirus 7 or aotine herpesviruses 1 and 3. Further, the herpesviruses may be selected from γ -herpesviruses, e.g. from

20 lymphocryptoviruses such as Epstein-Barr virus, cercopithecine herpesvirus 2 or porcine herpesvirus 1, or from rhadinoviruses such as human

herpesvirus 8, ateline herpesvirus 2 or saimiriine herpesvirus 1, or preferably, the virus is selected from human herpesvirus 1 (HSV-1), varicella

zoster virus (VZV) or human cytomegalovirus (HCMV).

25

Further, it was surprisingly found that the compounds are active against Foscarnet-resistant HCMV strains. Thus, they are likely to have a mode of

action which is different from the HCMV drugs on the market, making them valuable tools for combination therapy approaches, e.g. for combination

30 therapies together with viral DNA polymerase inhibitors. Furthermore, it was found that the compounds are potent inhibitors of ganciclovir-resistant virus strains, particularly ganciclovir-resistant HCMV strains.

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The compounds of the formula I are suitable for the manufacture of an agent for the prophylaxis and/or treatment of a viral infection. This prophylaxis or treatment comprises administering a pharmaceutical composition containing as an active agent a pharmaceutically effective amount of a compound of the general formula I to a subject, preferably a human, in need thereof, e.g. a subject suffering from a herpesvirus infection or a subject which is in need of a prophylactic administration to avoid the outbreak of a herpesvirus infection. The pharmaceutical composition may contain suitable diluents, carriers and auxiliary agents. Further, the composition may also contain other pharmaceutically active agents, e.g. antiviral agents. The pharmaceutical composition may be suitable for oral, parenteral, e.g. intradermal, intravenous or intramuscular, rectal, nasal and topical applications. The composition may be an injectable solution, ointment, cream or spray. Further, the composition may have retardation properties, i.e. showing a delayed release of the active agent.

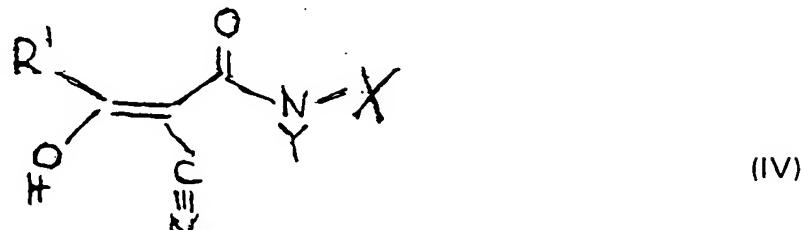
The dosage of the active agent depends on the specific compound being administered, the type and the severity of the viral infection. For example, a dosage from 0.01 mg to 100 mg per day and per kg/body weight for the active agent is suitable.

The efficacy of the compounds of formulae (Ia), (Ib), (Ic) and (Id) against enzymes from pathogens, e.g. viral kinases, may be determined in an in vitro enzyme inhibition test. On the other hand, the anti-viral effect can also be determined directly in a cell culture assay. A preferred and new cell culture assay is described in Example 2.2 and comprises the use of cells infected with a recombinant human cytomegalovirus carrying a reporter gene, e.g. the GFP-gene. The reporter gene is preferably inserted into the viral genome in a manner that viral replication is still possible. For example, the reporter gene may be inserted into the HCMV gene region encoding the open reading frames US9 and US10.

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A further embodiment of the present invention refers to the use of compounds of the general formula (IV):

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wherein X and Y are defined as for the compounds (Ia-d) and R' is selected from C₁-C₄ alkyl and C₃-C₄ cycloalkyl, particularly CH₃, for the preparation of an agent against ganciclovir-resistant virus strains. The compounds may be present in the form of a physiologically acceptable salt, e.g. an alkali metal, ammonium or substituted ammonium salt, particularly the sodium salt or the salt of a basic amino acid such as lysine. The compounds of formula (IV) may be classified as leflunomides. Surprisingly, these leflunomides are potent inhibitors of ganciclovir-resistant virus strains, particularly ganciclovir-resistant herpes virus strains and more particularly ganciclovir-resistant HCMV strains.

15

The compounds of the formula (IV) are suitable for the manufacture of an agent for the prophylaxis and/or treatment of a viral infection. This prophylaxis or treatment may be carried out as described above for compounds of the formulae (Ia) and (Ib).

Furthermore, the invention is to be explained by the following figures and examples.

25

Fig. 1 is a list of compounds of the present invention having anti-viral activity.

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Fig. 2 is a diagram showing the efficacy of compounds of the present invention against a ganciclovir-resistant virus strain.

Example 1

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Synthesis of substituted anilides of heterocyclic carboxylic acids

A general procedure is given here suitable for acylation of substituted anilines with heterocyclic carboxylic acids. The aromatic rings can be mono, di, tri, or tetrasubstituted with e.g. alkyl, substituted alkyl, aryl, hydroxy, halogen, nitro, carboxy, thiomethyl groups. The heterocyclic ring can be mono, di, tri, or tetrasubstituted with e.g. alkyl, substituted alkyl, aryl, hydroxy, oxo, halogen, nitro, thiomethyl groups.

15 **General procedure**

1-10 mmol aniline was dissolved in dry 10-30 mol of dry pyridine and the solution was cooled to -20°C. 0.5 eq. of phosphorous trichloride was dropped in with vigorous stirring while the temperature was kept under -15°C. After 10-15 min the carboxylic acid was added slowly with stirring. The reaction mixture was allowed to warm to room temperature and stirred overnight, then warmed to 40°C until 30 min. Pyridine was removed in vacuum and the residue was taken up in 1N hydrochloric acid and ethyl acetate. The organic phase was washed with 5% hydrocarbonate solution and distilled water. After drying over sodium sulfate the solvent was evaporated and the residue purified via filtrating through silica layer, or chromatographed or recrystallized, if necessary.

All reagents and intermediates were purchased from Sigma-Aldrich.

30

For example, the following compounds were obtained:

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(1) Thiophene-2-carboxylic acid (3-trifluoromethyl phenyl)amide M.p.: 151-152°C R_f:0.73 (Ethylacetate:Hexane = 1:1)

5 (2) Furane-2-carboxylic acid (3-trifluoromethyl phenyl)amide M.p.: 108-109°C R_f:0.73 (Ethylacetate:Hexane = 1:1)

(3) Thiophene-2-carboxylic acid (3,5 bis-(trifluoromethyl) phenyl)amide M.p.: 148°C R_f:0.8 (Ethylacetate:Hexane = 1:1)

10 (4) Furane-2-carboxylic acid (4-trifluoromethyl phenyl)amide M.p.: 108-109°C R_f:0.73 (Ethylacetate:Hexane)

(5) Furane-2-carboxylic acid (3,5 bis-(trifluoromethyl) phenyl)amide M.p.: 124-125°C R_f:0.85 (Ethylacetate)

15 (6) 2-Cyano-3-hydroxy-crotonic acid (4-trifluoromethyl phenyl) amide

According to the general procedure as described above, further compounds of the formula I can be prepared, e.g. by using corresponding pyrrol, 20 pyrrolidine or oxo-pyrrolidine carboxylic acids as starting materials. A list of such compounds is shown in Fig. 1.

Example 2

25 Anti-viral activity of compounds according to formulae (Ia) and (Ib)

2.1 DHODH assay

The cDNA for human DHODH (accession number M94065) was obtained 30 via PCR with EST clone AA173225 (from RZPD in Berlin) and oligonucleotides 5'-CTG AAT TCA AAT TAC CGT GGA GAC ACC TGC AAA AGC GGG CCC AG-3' (this oligo provides the coding sequence for the

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first 5 amino acids missing in the EST clone) and 5'-AGC TCG AGT CAC CTC CGA TGA TCT GCT CCA AT-3'. The PCR product was subcloned with EcoRI and Xhol into vector pGEX 5x1 (Pharmacia, Gene (1988) 67:31) for expression as a GST-fusion protein in bacteria. Recombinant GST-DHODH was purified on glutathione-sepharose (method described in Gene (1988) 67:31) and dialyzed against 20 mM Tris pH 7.5, 0.1% Triton X100. The assay was basically performed as described by Copeland et al. ("NBT assay", Arch.Biochem.Biophys.323:79, 1995) with slight modifications: 20 μ l of a 5x ubiquinone mixture (1 mM Ubiquinone Q10, 0.5% Triton X100, 10 500 mM Tris pH 7.5), 2 μ l of 10 mM NBT (nitroblue tetrazolium), 2-10 μ l of DHODH-GST (depending on the batch of 1-liter bacterial preparation), 91- 15 83 μ l of water and 5 μ l of compound or DMSO were added to each well of a 96-well microtiter plate. After mixing and incubation of the reactions for 5 min at 37°C, 5 μ l of 10 mM L-dihydroorotate (or 5 μ l of water in case of the background controls) were added to start the reactions. After incubation at 37°C for 60 to 90 min, ODs were read at 595 nm with a Biorad microplate reader. Readings were in duplicate, controls without L-dihydroorotate were performed as single measurements. Activity was normalized with values from DMSO controls.

20

2.2 Virus Replication Assay

Cell culture and virus

Primary human foreskin fibroblasts (HFF) were cultivated in MEM containing 25 5% (v/v) fetal calf serum. Infection analysis was restricted to cell passage numbers below twenty. Human cytomegalovirus strain AD169 (ATCC) was grown in HFF cells and quantitated for infectivity by the plaque reduction assay. Aliquots were stored at -80°C.

30 **Construction of recombinant cytomegalovirus**

For construction of a recombination vector, two linker sequences were inserted into the pBlueScribe vector pBS+ (Stratagene): the first contained

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restriction sites for NheI, SpeI, PacI and BglII followed by a loxP sequence (ATAACTTCGTATAGCATACATTATACGAAGTTAT) and was introduced into PstI/XbaI sites of the vector; the second contained another loxP sequence followed by restriction sites HpaI, ClaI and PmeI and was 5 introduced into BamHI/Asp718 sites. A gene cassette comprising of a "humanized" version of the ORF coding for GFP (gfp-h) under the control of the HCMV enhancer/promoter and the Ptk/PY441 enhancer-driven neoR selection marker was excised from plasmid pUF5 (Zolotukhin et al., 1996, J.Virol.70, 4646-4654) and inserted into the recombination vector via BglII 10 sites.

At the 5' and 3'-positions of this loxP-flanked gene cassette, two HCMV sequences with homology to the gene region containing the open reading frames US9 and US10 were inserted. For this, viral sequences were 15 amplified from template pCM49 (Fleckenstein et al., 1982, Gene 18, 39-46) via PCR in a 35-cycle program (denaturation 45 sec at 95°C, annealing 45 sec at 55°C and elongation 2 min at 72°C) by the use of Vent DNA polymerase (New England Biolabs). A US10-specific sequence of 1983 bp in length was generated using primers US10[200900]SpeI 20 (GCTCACTAGTGGCCTAGCCTGGCTCATGGCC) and US10[198918]PacI (GTCCTTAATTAAGACGTGGTTGTGGTCACCGAA) and inserted at the vector 5' cloning position via SpeI/PacI restriction sites (see bold-print). A US9-specific sequence of 2010 bp was generated using primers US9- 3'PmeI (CTCGGTTAAACGACGTGAGGCGCTCCGTCACC) and US-5' ClaI 25 (TTGCATCGATAACGGTGTGAGATACCAACGATG) inserted at the vector 3' cloning position via PmeI/ClaI restriction sites.

The resulting construct pHM673 was linearized by the use of restriction enzyme NheI and transfected into HEF cells via the electroporation method 30 using a Gene Pulser (Bio-rad; 280 V, 960 µF, 400 Ω). After 24 h of cultivation, cells were used for infection with 1 PFU/ml of HCMV strain AD169. Selection with 200 µg/ml G418 was started 24 h post infection.

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Following 3 weeks of passage in the presence of G418, GFP fluorescence could be detected in most of the infected cells. Plaque assays were performed with infectious culture supernatant on HFF cells and single virus plaques were grown by transfer to fresh HFF cells cultured in 48-well plates. DNA was isolated from cells of 32 fluorescence-positive wells and confirmed for the presence of recombinant virus by PCR. For this, primers 5 US9[198789] (TGACGCGAGTATTACGTGTC) and US10[199100] (CTCCTCCTGATATGCGGTT) were used resulting in an amplification product of 312 bp for wild-type AD169 virus and approximately 3.5 kb for 10 recombinant virus.

Plaque assay

HFF cells were cultivated in 12-well plates to 90-100% confluency and 15 used for infection with dilutions of virus-positive cell culture supernatants. Virus inoculation was performed for 90 min at 37°C under occasional shaking before virus was removed and the cell layers were rinsed with PBS. Overlays of MEM 5% (v/v) fetal calf serum and 0.3% (w/v) agarose were added to each well and all samples were incubated at 37°C in a 5% CO₂ 20 atmosphere for approximately 12 days. Finally, overlays were removed and the formation of foci was visualized by staining with 1% crystal violet in 20% ethanol for 1 min. After repeated rinsing with PBS, plates were air-dried at room temperature and plaque numbers were counted with a light microscope. For the recombinant AD169-GFP virus, quantification of plaque 25 assays could also be performed without crystal violet staining by a direct counting of the amount of green fluorescent plaques using fluorescence microscopy.

Antiviral compounds

30 The reference compounds used for antiviral studies, ganciclovir (GCV, Cymeven), foscarnet sodium (FOS, Foscavir) and cidofovir (CDV, Vistide) were purchased from Syntex Arzneimittel (Aachen, Germany), Sigma-

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Aldrich (Germany) and Pharmacia & Upjohn S.A. (Luxembourg), respectively. Stocks were prepared in aqueous solution and stored at -20°C. The test compounds were dissolved in DMSO and aliquots were stored at -20°C.

5

GFP infection assay

HFF cells were cultivated in 12-well plates to 90-100% confluence and used for infection with 0,5xTCID₅₀ of AD169-GFP virus. Virus inoculation was performed for 90 min at 37°C with occasional shaking before virus 10 was removed and the cell layers were rinsed with PBS. Infected cell layers were incubated with 2 ml of MEM containing 5% (v/v) fetal calf serum and optionally of the respective test substances or DMSO as control. Infected cells were incubated at 37°C in a 5% CO₂ atmosphere for 7 days and harvested by trypsinization and centrifugation. 200 µl of lysis buffer (25 mM 15 Tris pH 7.8, 2 mM DTT, 2 mM trans-1,2-diaminocyclohexane-N,N,N',N'-tetraacetic acid, 1% Triton X-100, 10% glycerol) was added to each cell pellet and lysis was achieved by incubation for 10 min at 37°C followed by a 30-min incubation at room temperature on a shaker. Lysates were centrifuged for 5 min at 15.000 rpm in an Eppendorf centrifuge to remove 20 cell debris. Supernatants were transferred to an opaque 96-well plate for automated measuring of GFP signals in a Victor 1420 Multilabel Counter (Wallac). GFP units were converted to percent inhibition values relative to DMSO controls (set at 100% GFP expression).

25 **Indirect immunofluorescence analysis**

Cells were either grown on Lab-Tek Permanox slides (Nunc) or harvested from 6-well plates, spotted onto glass slides with marked rings (Medco) and fixed by a 15-min treatment with 3% formaldehyde in PBS followed by permeabilization for 15 min in 0.1% Triton X-100 in PBS at room 30 temperature. Blocking was achieved by incubation with Cohn Fraction II/III of human gamma-globulin (Sigma; 2 mg/ml) for 30 min at 37°C. The IE1/IE2-specific primary antibody MA810 (Chemicon International, Inc. CA,

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USA; dilution 1:10.000) was incubated for 90 min, the secondary antibody (tetramethyl rhodamine [TRITC]-coupled anti-mouse antibody, Dianova, dilution 1:100) for 45 min at 37°C before analysis by fluorescence microscopy. In addition to indirect TRITC staining of IE1/IE2 proteins, GFP 5 signals could be detected directly via the fluorescence isothiocyanate (FITC) channel. Nuclear counterstaining was carried out using Vectashield mounting medium including DAPI (Vector Laboratories, Burlingame, CA).

2.3 Results

10

First, the ability of the compounds to inhibit the de novo pathway of pyrimidine biosynthesis, possibly leading to antiproliferative and immunosuppressive effects *in vivo*, was tested by determining DHODH activity *in vitro* in the presence of 20 μ M of the test compounds. The 15 maximum inhibition of DHODH was 26%, whereas the active metabolite of Leflunomide inhibited at least 80% of DHODH activity under test conditions. Compounds (1) and (3), two compounds with a thiophene ring, were very effective in inhibiting HCMV replication at concentrations of 20 and 100 μ M, while (2) which is identical to (1), except that it contains a furane ring 20 instead of a thiophene ring, showed slightly lower reduction at 20 μ M concentration. Compounds (4) and (5) inhibited HCMV significantly at the higher, but not at the lower concentration.

25 Table 1 shows the compounds and lists which percentage of DHODH activity is inhibited at a concentration of 20 μ M. Values are averages of duplicate measurements from two different experiments. Percentage of HCMV inhibition at 20 μ M and 100 μ M (two different experiments) of the substances are also shown. Values are averages of duplicate measurements.

30

Table 1 -

	DHOD inhibition [%]	HCMV inhibition [%]	
Compound	20 μ M	100 μ M	20 μ M
(1)	6 \pm 10	99 \pm 0	90 \pm 4
(2)	5 \pm 4	96 \pm 1	73 \pm 10
(3)	11 \pm 9	100 \pm 0	99 \pm 0
(4)	12 \pm 9	92 \pm 1	9 \pm 26
(5)	15 \pm 11	98 \pm 0	39 \pm 12

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The compounds shown in Fig. 1 also have significant anti-viral activity in the assay system as described above.

Example 3

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Anti-viral activity against ganciclovir-resistant virus strains

3.1 Isolation of Drug-Resistant Virus

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A series of laboratory variants of AD169-GFP virus with resistance against ganciclovir (GCV) was generated. HFF cells were infected in 12-well plates with MOI 0.002 and incubated with 1 μ M of GCV. GFP-expression in infected cells was monitored microscopically and the supernatants of positive wells were transferred to fresh cells weekly. Thereby GCV concentrations were increased stepwise (1- μ M increase in each step) up to the point where the total virus replication became critical and resistant virus grew out in individual wells. Using supernatants of these wells, two rounds of plaque purifications were performed on HFF cells. Finally, GCV-resistant

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- 18 -

viral clones (e.g. AD169-GFP314) were isolated which were able to replicate in the presence of 10 μ M of GCV.

3.2 Virus Replication Assay

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Plaque purification and plaque reduction assay

HFF cells were cultivated in 12-well plates to 90-100% confluence and used for infection with dilutions of virus stocks (i.e. AD169-GFP or AD169-GFP314). Virus inoculation was performed for 90 minutes at 37°C under occasional shaking before virus was removed and the cell layers were rinsed with PBS. Overlays of MEM containing 5% (v/v) fetal calf serum and 0.3% (w/v) agarose were added to each well. The plates were incubated at 37°C in a 5% CO₂ atmosphere for 8-12 d. For plaque purification of GFP-expressing viruses, plates were used for fluorescence microscopy, GFP-positive plaques were picked from the overlays and transferred to fresh cells for virus multiplication. For plaque reduction assays, antiviral compounds were incubated in the overlays after infection. Overlays were removed and plaque formation was visualized by staining with 1% cristal violet in 20% ethanol for 1 min. After repeated rinsing with PBS, plates were air-dried at room temperature and plaque numbers were counted with a light microscope. For the GFP-expressing recombinant viruses, quantification of plaque reduction assays could be performed alternatively, without cristal violet staining, by a direct counting of the numbers of green fluorescent plaques using fluorescence microscopy.

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- 19 -

one of the respective test compounds. Infected cells were incubated at 37°C in a 5% CO₂ atmosphere for 7 d. For lysis, 200 µl of lysis buffer (25 mM Tris pH 7.8 2 mM DTT, 2 mM trans-1,2-diaminocyclohexane-N,N,N',N'-tetraacetic acid, 1% Triton X-100, 10% glycerol) was added to each well and incubated for 10 min at 37°C followed by a 30-min incubation at room temperature on a shaker. Lysates were centrifuged for 5 min at 15,000 rpm in an Eppendorf centrifuge to remove cell debris. 100 µl of the supernatants were transferred to an opaque 96-well plate for automated measuring of GFP signals in a Victor 1420 Multilabel Counter (Wallac).

10

3.3 Results

GCV-resistant HCMV is sensitive to compound (1) or compound (6).

15 HFF cells were cultivated in 12-well plates and infected with the indicated concentrations of GFP-expressing variants of HCMV, AD169-GFP (parental) or AD169-GFP314 (GCV-resistant). Immediately after the infection, the following chemical compounds were added to the culture media: DMSO 0.07% (A), 0.14% (B), ganciclovir (GCV) 10 µM (A and B), compound (6) 20 35 µM (A) and 70 µM (B), and compound (1) 35 µM (A) and 70 µM (B). Seven days postinfection infected cells were harvested and used for GFP quantification. The results for parental virus AD 169-GFP and a GCV-resistant virus mutant (AD169-GFP314) are shown in Fig. 2a and 2b, respectively.

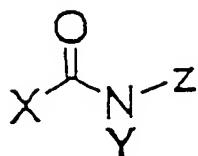
- 20 -

Claims

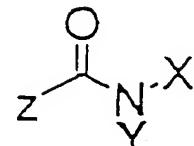
1. Use of compounds of the general formulae (Ia), (Ib), (Ic) or (Id):

5

(Ia)

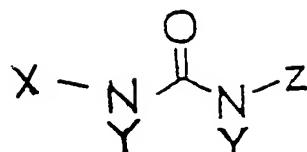


(Ib)

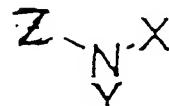


15

(Ic)



(Id)



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wherein X and Z are substituents comprising an aromatic or heteroaromatic ring system and Y is hydrogen or C₁-C₄ alkyl as well as the pharmaco logically acceptable salts thereof, for the preparation of an agent against infectious diseases.

2. Use of claim 1,

wherein X and/or Z is an aromatic radical which is unsubstituted or which carries at least one substituent which may be selected from hydroxy, cyano, nitro, halo, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkoxy, C₁-C₄ alkylthio, C₁-C₄ haloalkylthio, carboxy, carboxy-C₁-C₄-alkyl, carboxy-aryl or -heteroaryl,

- 21 -

aminocarbonyl - C₁-C₄-alkyl, amino carbonyl - aryl or - heteroaryl, aryl and heteroaryl.

3. Use according to claim 1 or 2,

5 wherein X and/or Z is a phenyl radical which may carry up to 4 substituents.

4. Use according to claim 1 or 2,

wherein X and/or Z carries at least one C₁-C₃ haloalkyl substituent.

10

5. Use according to claim 4,

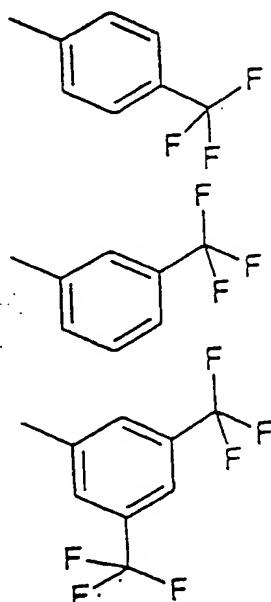
wherein X and/or Z carries at least one substituent selected from -CF₃, -CHF₂ and -CH₂F.

15

6. Use according to claim 5,

wherein X is selected from radicals represented by the formulae (IIa-c):

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- 22 -

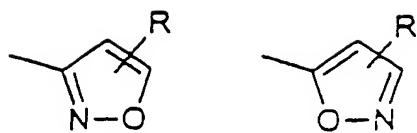
7. Use according to any of the claims 1 - 6,
wherein X and/or Z comprises a heterocyclic ring.
8. Use according to any one of claims 1-5,
5 wherein X and/or Z comprises a five-membered heterocyclic ring, a
six-membered heterocyclic ring or a bi- or polyheterocyclic ring.
9. Use according to claim 8,
10 wherein X and/or Z comprises a five-membered heterocyclic ring
selected from pyrrole, pyrazole, imidazole, 1,2,3-triazole, tetrazole,
oxazole, isoxazole, thiazole, isothiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole, thiophene, furan, indole and 3-thiaindole.
10. Use according to claim 9,
15 wherein X and/or Z is selected from radicals represented by the
formulae (IIIa-k):

20

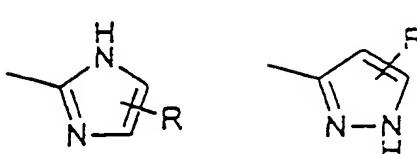
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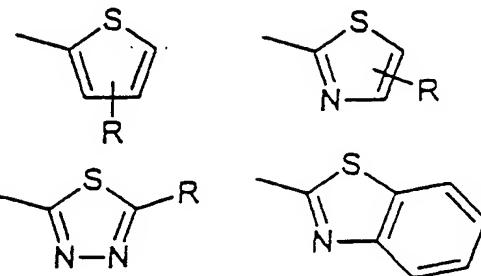
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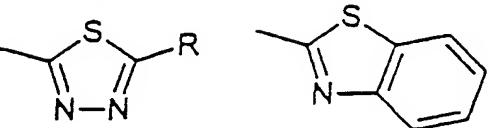
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15



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wherein R is at least one substituent selected from halo, C₁-C₃ alkyl and C₁-C₃ alkoxy.

25

11. Use according to any one of claims 1-10,
wherein compounds (Ia), (Ib), (Ic) and (Id) have a higher selectivity
for kinases from pathogens than for the cellular enzyme DHODH.
- 30 12. Use according to any one of claims 1-11 for the preparation of an
agent against viral infections.

- 24 -

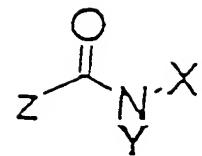
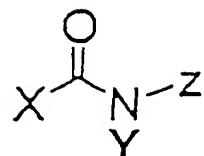
13. Use according to claim 12 for the preparation of an agent against infections by herpesviruses.
14. Use according to claim 13,
5 wherein the virus is selected from herpes simplex viruses, varicella viruses, cytomegaloviruses, muromegaloviruses, roseoloviruses, lymphocryptoviruses and rhadinoviruses.
15. Use according to claim 14,
10 wherein the virus is human cytomegalovirus (HCMV).
16. Use according to any one of claims 12-15,
wherein the virus is a Foscarnet- or ganciclovir-resistant virus strain.
- 15 17. Use according to any one of claims 1-16,
wherein the agent is suitable for oral, parenteral, rectal, nasal and topical application.
18. Use according to any one of claims 1-17,
20 wherein the dosage of the active agent is from 0.01 mg to 100 mg per day and per kg body weight.
19. Use according to any one of claims 1-18 as a combination therapy with viral DNA polymerase inhibitors.
- 25 20. A method for the treatment or prophylaxis of a herpesvirus infection comprising administering to a subject in need thereof a pharmaceutically effective amount of a compound according to general formulae (Ia), (Ib), (Ic) or (Id):

- 25 -

(Ia)

(Ib)

5

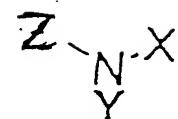
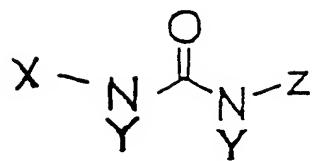


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(Ic)

(Id)

15



wherein X and Z are substituents comprising an aromatic or heteroaromatic ring system and Y is hydrogen or C₁-C₄ alkyl.

20

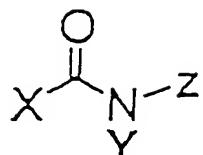
21. Compounds according to the general formulae (Ia), (Ib), (Ic) or (Id):

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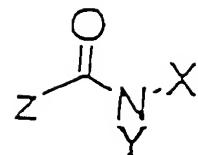
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- 26 -

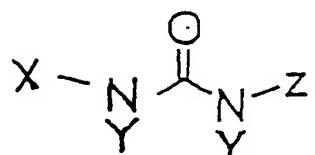
(Ia)



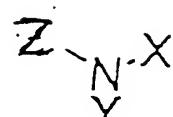
(Ib)



(Ic)



(Id)

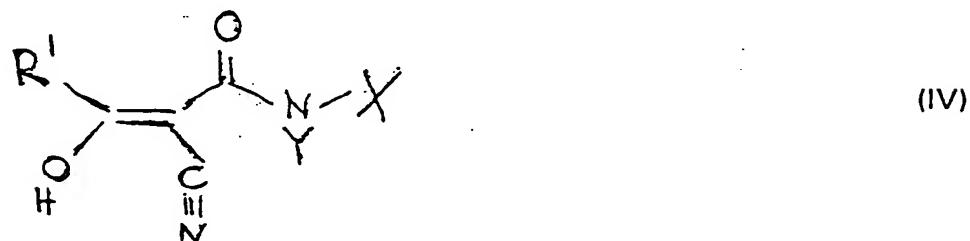


wherein X and Z are substituents comprising an aromatic or heteroaromatic ring system and Y is hydrogen or C₁-C₄ alkyl.

20

22. Use of compounds of the general formula (IV)

25



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wherein X and Y are defined as in any one of claims 1-7 and R' is C₁-C₄ alkyl or C₃-C₄ cycloalkyl as well as the pharmacologically

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acceptable salts thereof, for the preparation of an agent against ganciclovir-resistant virus strains.

23. Use according to claim 22,

5 wherein R' is CH₃.

24. Use according to claim 22 or 23,

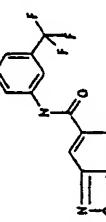
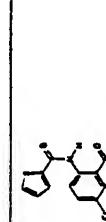
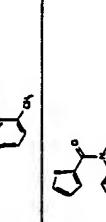
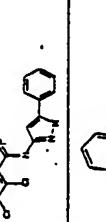
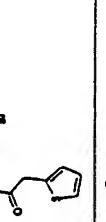
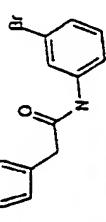
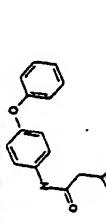
wherein the virus is human cytomegalovirus (HMCV).

-1/6-

Fig. 1

IUPAC NAME	STRUCTURE	MP	mol weigh	TLC Rf.
N-(5-Phenyl-1H-pyrazol-3-yl)-4-trifluoromethyl-benzamide		234 - 235	331,2999	0.59(B) 0.63(C)
4-Trifluoromethyl-N-(4-trifluoromethyl-phenyl)-benzamide		104 - 106	333,2355	0.79 (B) 0.51(C)
N-Quinolin-8-yl-trifluoromethyl-benzamide		75 - 78	316,2852	0.93(B) 0.51(C) 0.78(D)
N-(4-iodo-phenyl)-4-trifluoromethyl-benzamide		148 - 151	391,1335	0.82(B) 0.49(C) 0.84(D)
N-(3-(4-trifluoromethyl-benzoyl)-amino-phenyl)-4-trifluoromethyl-benzamide		187 - 189	452,3595	0.61(B) 0.53(D)
N-(4-Cyano-phenyl)-4-trifluoromethyl-benzamide		179 - 180	290,247	0.64(B) 0.51(D)
3,4-Dimethyl-thieno[2,3-b]thiophene-2-carboxylic acid[(bis-trifluoro-methyl)-phenyl]amide			423,4017	Maybridge KM 03523
Quinoline-6-carboxylic acid (3-trifluoromethyl-phenyl)-amide		118 - 120	316,2852	KM 10124
		157 - 160		

continued Fig. 1

Benz[1,2,5]oxadiazole-5-carboxylic acid > (3-trifluoromethyl-phenyl)-amide		180 - 183	307,234	Maybridge KM 09572
Thiophene-2-carboxylic acid[3,4-dichloro-2-(3,4-dimethyl-phenyl)]-amide		222 - 223	419,3327	0.61(A) 0.67(B) 0.42(D)
Thiophene-2-carboxylic acid[3,4-dichloro-2-(5-phenyl-1H-pyrazol-3-ylcarbamoyl)-phenyl]-amide		254 - 255	457,3413	0.35(A) 0.50(B) 0.06(D)
N-(3-Bromo-phenyl)-2-(thiophen-2-yl)-acetamide		98 - 100	296,1876	0.56(A) 0.54(B) 0.36(D)
N-(3-Bromo-phenyl)-2-(thiophen-3-yl)-acetamide		100 - 101	296,1876	0.55(A) 0.55(B) 0.34(D)
N-(4-Phenoxy-phenyl)-2-(thiophen-2-yl)-acetamide		98 - 101	309,3890	0.52(A) 0.58(B) 0.32(D)
Thiophene-3-carboxylic acid (4-bromo-phenyl)-amide		194 - 196	282,1605	0.56(A) 0.51(B) 0.38(D)
N-(4-Bromo-phenyl)-2-(thiophen-3-yl)-acetamide		150	296,1876	0.54(A) 0.51(B) 0.29(D)

continued Fig. 1

2-Thiophen-3-yl-N-(4-trifluoromethyl-phenyl)-acetamide		177 - 178	285,29	0.56(A) 0.50(B) 0.33(D)
Thiophene-2-carboxylic acid biphenyl-4-ylamide		203 - 204	279,3633	0.55(A) 0.55(B) 0.33(D)
N-(4-(4-trifluoromethyl-benzenesulfonyl)-amino-pyrimidin-2-yl)-4-trifluoromethyl-phenylsulfonamide		175 - 176	526,4391	0.36(A) 0.48(B) 0.03(D)
Thiophene-2-carboxylic acid (3-trifluoromethyl-phenyl)-amide		128 - 130	271,2629	0.58(A) 0.46(B) 0.30(D)
Thiophene-2-carboxylic acid (4-trifluoromethyl-phenyl)-amide		170 - 172	271,2629	0.53(B) 0.35(D)
Thiophene-2-carboxylic acid (bis-trifluoromethyl-phenyl)-amide		145 - 146	339,2612	0.63(A) 0.47(B) 0.34(D)
(Bis-trifluoromethyl-phenyl)-(dimethyl-1H-benzimidazol-2-yl)-amine		242 - 248	373,3037	0.57(A) 0.39(B) 0.14(D)
1-(Amino-dichloro-phenyl)-3-(bis-trifluoromethyl-phenyl)-urea		350	432,1549	0.56(A) 0.39(B) 0.10(D)

continued Fig. 1

(3,5-Bis(trifluoromethyl)-phenyl)-(4-methyl-1H-benzimidazol-2-yl)-amine		232 - 235	359,2766	0.56(A) 0.40(B) 0.13(D)
Thiophene-2-carboxylic acid [2-(benzothiazol-2-ylcarbamoyl)-3,4-dichloro-phenyl]-amide		274 - 275	448,3524	0.52(A) 0.65(B) 0.18(D)
Thiophene-2-carboxylic acid [5-chloro-2-(3,4-dimethyl-phenylcarbamoyl)-phenyl]-amide		196 - 197	384,8877	0.61(A) 0.60(B) 0.31(D)
Thiophene-2-carboxylic acid [2-(4-bromo-phenylcarbamoyl)-5-chloro-phenyl]-amide		236 - 237	435,7295	0.61(A) 0.59(B) 0.29(D)
Thiophene-2-carboxylic acid [2-(4-acetylamino-phenylcarbamoyl)-4-bromo-phenyl]-amide		281 - 288	458,3368	0.35(A) 0.44(B) 0.02(D)
Thiophene-2-carboxylic acid [2-(benzothiazol-2-ylcarbamoyl)-4-bromo-phenyl]-amide		252 - 255	458,3584	0.56(A) 0.65(B) 0.21(D)
Thiophene-2-carboxylic acid [2-(4-bromo-phenylcarbamoyl)-4-chloro-phenyl]-amide		218 - 220	435,7295	0.61(A) 0.63(B) 0.31(D)
Thiophene-2-carboxylic acid [2-(4-bromo-phenylcarbamoyl)-4-chloro-phenyl]-amide		257 - 260	413,9074	0.57(A) 0.65(B) 0.25(D)
Thiophene-2-carboxylic acid [2-(benzothiazol-2-ylcarbamoyl)-4-chloro-phenyl]-amide				

continued Fig. 1

Thiophene-2-carboxylic acid [4-chloro-2-(thiazol-2-ylcarbamoyl)-phenyl]-amide		190	363,8469	0.51(A) 0.60(B) 0.13(D)
Thiophene-2-carboxylic acid [2-(1H-benzoimidazol-2-ylcarbamoyl)-3-chloro-phenyl]-amide		237 - 238	396,8581	0.35(A) 0.49(B) 0.03(D)
Thiophene-2-carboxylic acid [2-(benzothiazol-2-ylcarbamoyl)-3-chloro-phenyl]-amide		239	413,9074	0.47(A) 0.59(B) 0.13(D)
Thiophene-3-carboxylic acid (3-trifluoromethyl-phenyl)-amide		119	271,2629	0.57(A) 0.47(B) 0.29(D)
3-Trifluoromethyl-N-(4-trifluoromethyl-phenyl)-benzamide		115 - 116	333,2355	0.62(A) 0.52(B) 0.34(D)
TLC conditions:				
A: Hexane: Acetone 1:1				
B: Benzene: Methanol 1:4				
C: Ethylacetate: Hexane 1:5				
D: Ethylacetate: Hexane 1:2				

Fig. 2

